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In the Claims:

Please amend the claims as shown below. This listing of claims will replace all prior versions and listings of the claims in this application.

- 1. (Currently amended) A method of regulating apoptosis in a cell, said method comprising targeting an abnormally alternatively spliced <u>Bcl-2</u> mRNA, an abnormally alternatively structured <u>Bcl-2</u> mRNA, or a product of either.
- 2. (Currently amended) A <u>The</u> method according to claim 1 further comprising targeting the junctions of the <u>Bcl-2</u> mRNA molecule that is abnormally spliced or abnormally structured.
- 3. (Currently amended) A <u>The</u> method according to claim 1 further comprising targeting a protein product following translation of the abnormally spliced or abnormally structured Bcl-2 mRNA.
- 4. (Currently amended) A <u>The</u> method according to <u>any of claimsclaim</u> 1 to 3 further comprising the selective silencing of abnormal splice variants of the Bcl-2 gene.
- 5. (Currently amended) A <u>The</u> method according to claim 4 further comprising the targeting of any of the abnormal splice variants selected from the group consisting of: Bcl-2 α -591, Bcl-2 α -588, Bcl-2 α -480, Bcl-2 α -633, Bcl-2 β -489, Bcl-2 β -474, Bcl-2 β -420 and/or Bcl-2 β -315.
- 6. (Currently amended) A <u>The</u> method according to claim 5 further comprising targeting of the mRNA sequence flanking the splice junction between nucleotides 111 and 241 of Bcl- 2α -591.
- 7. (Currently amended) A <u>The</u> method according to any of the preceding elaimsclaim 1 further comprising targeting an abnormally spliced <u>Bcl-2</u> mRNA

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or a product thereof, by introducing into a cell containing a <u>Bcl-2</u> gene which is abnormally spliced and which is to be targeted, an RNA construct having a nucleotide sequence which is homologous to mRNA within said cell wherein said mRNA includes genetic information of the <u>Bcl-2</u> gene element that is abnormally spliced.

- 8. (Currently amended) A <u>The</u> method according to claim 7 wherein the RNA construct is a small interfering dsRNA (siRNA).
- 9. (Currently amended) A <u>The</u> method according to claim 8 wherein the siRNA is up to 28 nucleotides long.
- 10. (Currently amended) A <u>The</u> method according to <u>any of claimsclaim</u> 1, to 6, further comprising targeting an abnormally spliced <u>Bcl-2</u> mRNA or a product thereof, by introducing into a cell containing a <u>Bcl-2</u> gene which is abnormally spliced and which is to be targeted, an agent selected from the group consisting of: small molecule or protein; polypeptide; peptide; aptamer; chemical; antibody; nucleic acid; polypeptide or nucleotide probe; anti-sense RNA; shRNA; miRNA; and Bcl-2 derived products including abnormal Bcl-2 splice variants which inhibit Bcl-2 either directly or indirectly; which agent interacts with or binds with the abnormally spliced <u>Bcl-2</u> mRNA or protein expressed by the abnormally spliced <u>Bcl-2</u> mRNA.
- 11. (Currently amended) A nucleotide construct with a nucleotide sequence which is at least 50% homologous to mRNA transcribed from an abnormally spliced Bcl-2 gene.
- 12. (Currently amended) A <u>The</u> nucleotide construct according to claim 11 wherein said construct comprises dsRNA.
- 13. (Currently amended) A <u>The</u> nucleotide construct according to claim 12 wherein the construct is 20 to 28 nucleotides long.

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- 14. (Currently amended) A <u>The</u> nucleotide construct according to claim 13 wherein the RNA construct is 21 to 22 nucleotides long.
- 15. (Currently amended) A nucleotide construct such as selected from the group consisting of: siRNA; anti-sense RNA; shRNA; and miRNA; as means for silencing the expression of an abnormally spliced <u>Bcl-2</u> gene for use as a medicament.
- 16. (Currently amended) An agent selected from the group consisting of: small molecule or protein; polypeptide; peptide; aptamer; chemical; antibody; nucleic acid; polypeptide; orand nucleotide probe; which agent interacts with or binds with a protein expressed by an abnormally spliced <u>Bcl-2 mRNA</u>, for use as a medicament.
- 17. (Currently amended) A <u>Use of a nucleotide construct such-asselected from the group consisting of:</u> siRNA; anti-sense RNA; shRNA; erand miRNA; and capable of silencing the expression of an abnormally spliced Bcl-2 gene for the manufacture of a medicament for the treatment of cancerous cell growth.
- 18. (Currently amended) AnUse of an agent selected from the group consisting of: small molecule or protein; polypeptide; peptide; aptamer; chemical; antibody; nucleic acid; polypeptide; erand nucleotide probe; which agent interacts with or binds with a protein expressed by an abnormally spliced Bcl-2 mRNA, for the manufacture of a medicament for the treatment of cancerous cell growth.
- 19. (Currently amended) A pharmaceutical composition comprising a nucleotide construct such as capable of silencing the expression of an abnormally spliced Bcl-2 gene and selected from the group consisting of: siRNA; anti-sense RNA; shRNA; erand miRNA; and a pharmaceutically acceptable diluent or carrier.

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- 20. (Currently amended) A pharmaceutical composition comprising an agent selected from the group consisting of: small molecule or protein; polypeptide; peptide; aptamer; chemical; antibody; nucleic acid; polypeptide; and nucleotide probe; which agent interacts with or binds with a protein expressed by an abnormally spliced <u>Bcl-2</u> mRNA, and a pharmaceutically acceptable diluent or carrier.
- 21. (Currently amended) Use of a DNA or RNA expression vector as a delivery means for a molecule which is used in the targeting of an abnormally spliced <u>Bcl-2</u> mRNA or a product thereof.
- 22. (Currently amended) A DNA or RNA expression vector comprising an expression cassette including the nucleotide sequence selected from the group consisting of; of:
- a) the nucleic acid sequence of an abnormally spliced gene element as shown in Fig 1;
- b) a nucleic acid molecule which <u>has at least 50% homology to and</u> hybridizes to the <u>a</u> nucleic acid sequence of (a); <u>and</u>
- c) a nucleic acid molecule which has a nucleic acid sequence which is degenerate because of the genetic code to the sequences in a) and b) and any sequence which is <u>at least 50%</u> complimentary to any of the above sequences;

wherein the expression cassette is transcriptionally linked to a promoter sequence.